

Listing of Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Currently amended) A process for manufacture of long circulating non-pegylated liposomes comprising: [[:]]

forming a lipid film by evaporating a solvent from a lipid solution comprising one or more phospholipids, a sterol and a solvent; hydrating the lipid film with an aqueous hydration media to form non pegylated liposomes;

wherein the amount of aqueous hydration media used is in the range of 10 to 35 ml for each mmole of phospholipid present in the lipid solution, and wherein the aqueous hydration media comprises ammonium sulfate and sucrose.

2. (Original) The process of claim 1 wherein the amount of aqueous hydration media used is 30 ml for each mmole of phospholipid in the lipid solution.

3. (Original) The process of manufacture of non-pegylated liposomes of claim 1 further comprising loading the liposomes with a therapeutic or diagnostic agent.

4. (Original) The process of claim 3, wherein the therapeutic agent is an antineoplastic agent.

5. (Original) The process of claim 4, wherein the antineoplastic agent is selected from the group consisting of Doxorubicin hydrochloride, Daunorubicin hydrochloride, and Epirubicin hydrochloride.

6. (Original) The process of claim 5, wherein the antineoplastic agent is Doxorubicin hydrochloride.

7. (Original) The process of claim 1, wherein the molar ratio of phospholipid to sterol is

from about 1:0.1-1:2.

8. (Original) The process of claim 7, wherein the wherein the molar ratio of phospholipid to sterol is from about 1:0.7.

9. (Canceled).

10. (Original) The process of claim [[9]] 1, wherein the concentration of ammonium sulfate in aqueous hydration media is not less than 125 mmoles/liter.

11. (Original) The process of claim 1, wherein the phospholipid has a phase transition temperature of 40°C. to 60°C.

12. (Original) The process of claim 11, wherein the phospholipid has a minimum of sixteen carbons fatty acid chain.

13. (Original) The process of claim 12, wherein the phospholipid is selected from the group consisting of Distearoyl phosphatidylcholine (DSPC), Dipalmitoyl phosphatidylcholine (DPPC), Hydrogenated soya phosphatidylcholine (HSPC) and derivatives of such phospholipids.

14. (Original) The process of claim 13, wherein the phospholipid is distearoyl phosphatidylcholine (DSPC) and wherein the sterol is cholesterol.

15. (Original) The process of claim 1, wherein the non-pegylated liposomes are successively extruded through series of filters having pore sizes from 0.4 μm to 0.05 μm for sizing.

16. (Original) A liposome manufactured by the process of claim 1.

17. (Original) The liposome of claim 16, wherein the phospholipid comprises distearoyl phosphatidylcholine (DSPC) and the sterol comprises cholesterol.

18. (Original) The liposome of claim 16, wherein the non-pegylated liposome further comprises a therapeutic or diagnostic agent.
19. (Original) The liposome of claim 18, wherein said therapeutic agent comprises an antineoplastic agent.
20. (Original) The liposome of claim 19, wherein the antineoplastic agent is selected from the group consisting of Doxorubicin hydrochloride, Daunorubicin hydrochloride, and Epirubicin hydrochloride.
21. (Original) The liposome of claim 20, wherein the antineoplastic agent is Doxorubicin hydrochloride.
22. (Original) The liposome of claim 16, wherein the average size of liposome is 0.06 μm to 0.16 μm in diameter.
- 23.-60 (Canceled)
61. (new) The process of claim 1 further comprising the step of removing the solvent before or after hydrating the lipid film.
62. (new) The process of claim , further comprising removing the solvent before after hydrating the lipid film; wherein the amount of the aqueous hydration media used is in the range of 10 to 35 ml for each mmole of phospholipid present in the lipid solution; sizing the non-pegylated liposomes to about 0.06 μm to 0.1 μm to form a liposomal composition; removing extra-liposomal hydration salt from the liposomal composition using sucrose-histidine buffer solution to form non-pegylated sized liposomes.